## **Guidelines Regarding Changes to Animal Study Proposals**

The Animal Welfare Act Regulations and the Public Health Service Policy on Humane Care and Use of Laboratory Animals (PHS Policy) require that Animal Care and Use Committees (ACUCs) review and approve changes to previously approved animal activities described in an Animal Study Proposal (ASP).

In accordance with Office of Laboratory Animal Welfare (OLAW) Guidance on Significant Changes to Animal Activities<sup>1</sup>, the NIH intramural research program Animal Research Advisory Committee (ARAC) recognizes that the individual Institute/Center (IC) ACUCs have the option to establish and approve IC-specific policies and guidance documents for the review and approval of proposed changes to ASPs. Without an IC-specific policy, approval of significant changes can only be granted after full committee review (FCR) or designated member review (DMR).

Several categories of proposed changes are outlined in OLAW guidance. These categories allow IC-ACUCs some discretion to use internal policies and guidance documents to define what constitutes a significant change or to establish a mechanism for determining significance on a case-by-case basis in accordance with the PHS Policy.

OLAW defines the following categories of proposed changes:

- 1. Significant changes that <u>must</u> be approved by FCR or DMR:
  - a) Changes from non-survival to survival surgery;
  - b) Changes resulting in greater pain, distress, or degree of invasiveness;
  - c) Changes in housing to a location not part of the program;
  - d) Changes in species;
  - e) Changes in study objectives;
  - f) Changes in Principal Investigator (PI); and
  - g) Changes that impact personnel safety.
- **2. Significant changes** that <u>may</u> be handled *administratively* according to an IC-ACUC-approved policy and in consultation with an IC-ACUC-authorized veterinarian:
  - a) Changes in anesthesia, analgesia, sedation, or experimental substances;
  - b) Changes in euthanasia to any method approved in the AVMA Guidelines; and
  - c) Changes in duration, frequency, type, or number of procedures performed on an animal.

These significant changes may be handled administratively through the optional process of veterinary verification and consultation (VVC). VVC allows an IC-ACUC-authorized veterinarian to verify that a proposed significant change is within the scope of an IC-ACUC-approved policy,

consistent with an IC-ACUC-approved guidance document, and appropriate for the animals and circumstance. The veterinarian is not conducting DMR but is serving as a subject matter expert and may refer any significant change to the IC-ACUC for FCR or DMR for any reason. Any significant change that does not meet the parameters of the IC-ACUC-approved policy must be referred to the IC-ACUC for FCR or DMR.

This process is contingent upon an IC-ACUC-approved policy that defines the scope of each type of significant change by listing a concurrent IC-ACUC-approved guidance document (SOP, formulary, guideline, etc.) upon which the change can be verified by the IC-ACUC-authorized veterinarian. The consultation with the veterinarian must be documented. The approved change must be documented in the ASP; must be sent to the appropriate animal facility staff; and the IC-ACUC should be kept informed of changes made through this process.

**3. Significant changes** that <u>may</u> be handled *administratively* according to an IC-ACUC-approved policy without additional consultation:

Increase in previously approved animal numbers.

This process is contingent upon an IC-ACUC-approved policy that defines the scope of changes in the number of animals permitted. No additional consultation is required. The approved change must be documented in the ASP, must be sent to the appropriate animal facility staff, and the IC-ACUC should be kept informed of changes made through this process.

Note: PIs may use fewer animals than approved. This does not require IC-ACUC-approval, notification, consultation, or administrative handling.

- **4. Other changes** that <u>may</u> be handled *administratively* without an IC-ACUC-approved policy, consultation, or notification:
  - a) Correction of typographical errors;
  - b) Correction of grammar;
  - c) Contact information updates; and
  - d) Change in personnel, other than the PI (provided all criteria required by the IC-ACUC is met).

This process does not require an IC-ACUC-approved policy or additional consultation. The approved change must be documented in the ASP and must be sent to the appropriate animal facility staff. For personnel changes, there must be an administrative review to ensure that all personnel are appropriately identified; adequately trained and qualified; enrolled in occupational health and safety programs; and meet other criteria as required by the IC-ACUC.

The IC-ACUC policies and guidance documents must be reviewed by the IC-ACUC at intervals of no less than once every three years to ensure they are appropriate and accurate. It is the IC-ACUC's responsibility to clearly define and communicate its policy for determining significance to PIs.

## Reference:

NIH OLAW Notice Number: NOT-OD-14-126, August 26, 2014 – Guidance on Significant Changes to Animal Activities: http://grants.nih.gov/grants/guide/notice-files/NOT-OD-14-126.html

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